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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.
10/533,734	05/19/2006	Peter Franz Ertl	PG5023	5039
20462 7590 09/17/2007 SMITHKLINE BEECHAM CORPORATION CORPORATE INTELLECTUAL PROPERTY-US, UW2220 P. O. BOX 1539 KING OF PRUSSIA, PA 19406-0939			EXAMINER KINSEY, NICOLE	
			ART UNIT 1648	PAPER NUMBER
			NOTIFICATION DATE 09/17/2007	DELIVERY MODE ELECTRONIC

Please find below and/or attached an Office communication concerning this application or proceeding.

The time period for reply, if any, is set in the attached communication.

Notice of the Office communication was sent electronically on above-indicated "Notification Date" to the following e-mail address(es):

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Office Action Summary

Application No.

10/533,734

Applicant(s)

ERTL, PETER FRANZ

Examiner

Nicole E. Kinsey, Ph.D.

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-- The MAILING DATE of this communication appears on the cover sheet with the correspondence address --

Period for Reply

A SHORTENED STATUTORY PERIOD FOR REPLY IS SET TO EXPIRE 3 MONTH(S) OR THIRTY (30) DAYS, WHICHEVER IS LONGER, FROM THE MAILING DATE OF THIS COMMUNICATION.

- Extensions of time may be available under the provisions of 37 CFR 1.136(a). In no event, however, may a reply be timely filed after SIX (6) MONTHS from the mailing date of this communication.
- If NO period for reply is specified above, the maximum statutory period will apply and will expire SIX (6) MONTHS from the mailing date of this communication.
- Failure to reply within the set or extended period for reply will, by statute, cause the application to become ABANDONED (35 U.S.C. § 133). Any reply received by the Office later than three months after the mailing date of this communication, even if timely filed, may reduce any earned patent term adjustment. See 37 CFR 1.704(b).

Status

- 1) ☒ Responsive to communication(s) filed on 28 June 2007.
- 2a) ☐ This action is **FINAL**. 2b) ☒ This action is non-final.
- 3) ☐ Since this application is in condition for allowance except for formal matters, prosecution as to the merits is closed in accordance with the practice under *Ex parte Quayle*, 1935 C.D. 11, 453 O.G. 213.

Disposition of Claims

- 4) ☒ Claim(s) 1-24, 28-31 and 36 is/are pending in the application.
- 4a) Of the above claim(s) _____ is/are withdrawn from consideration.
- 5) ☐ Claim(s) _____ is/are allowed.
- 6) ☒ Claim(s) 1-24, 28-31 and 36 is/are rejected.
- 7) ☐ Claim(s) _____ is/are objected to.
- 8) ☐ Claim(s) _____ are subject to restriction and/or election requirement.

Application Papers

- 9) ☒ The specification is objected to by the Examiner.
- 10) ☐ The drawing(s) filed on _____ is/are: a) ☐ accepted or b) ☐ objected to by the Examiner.
- Applicant may not request that any objection to the drawing(s) be held in abeyance. See 37 CFR 1.85(a).
- Replacement drawing sheet(s) including the correction is required if the drawing(s) is objected to. See 37 CFR 1.121(d).
- 11) ☐ The oath or declaration is objected to by the Examiner. Note the attached Office Action or form PTO-152.

Priority under 35 U.S.C. § 119

- 12) ☐ Acknowledgment is made of a claim for foreign priority under 35 U.S.C. § 119(a)-(d) or (f).
- a) ☐ All b) ☐ Some * c) ☐ None of:
1. ☐ Certified copies of the priority documents have been received.
2. ☐ Certified copies of the priority documents have been received in Application No. _____.
3. ☐ Copies of the certified copies of the priority documents have been received in this National Stage application from the International Bureau (PCT Rule 17.2(a)).

* See the attached detailed Office action for a list of the certified copies not received.

Attachment(s)

- 1) ☒ Notice of References Cited (PTO-892)
- 2) ☐ Notice of Draftsperson's Patent Drawing Review (PTO-948)
- 3) ☒ Information Disclosure Statement(s) (PTO/SB/08)
Paper No(s)/Mail Date 5/4/2005.
- 4) ☐ Interview Summary (PTO-413)
Paper No(s)/Mail Date. _____.
- 5) ☐ Notice of Informal Patent Application
- 6) ☐ Other: _____.

DETAILED ACTION

Applicant's election without traverse of Group I (claims 1-24, 28-31 and 36) in the reply filed on June 28, 2007 is acknowledged.

Specification

The title of the invention is not descriptive. A new title is required that is clearly indicative of the invention to which the claims are directed.

Applicants are required to update the first paragraph of the specification to provide the current continuing data (i.e., reference to PCT/EP03/12402) for the instant application.

The disclosure is objected to for not containing a brief description of the drawings section. According to 37 CFR § 1.74, when there are drawings, there shall be a brief description of the several views of the drawings and the detailed description of the invention shall refer to the different views by specifying the numbers of the figures, and to the different parts by use of reference letters or numerals (preferably the latter).

The disclosure is objected to because of the following informalities: Pages 45 and 46 contain amino acid sequences without sequence identifiers (SEQ ID NO:X).

Appropriate correction is required.

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Claim Rejections - 35 USC § 112

The following is a quotation of the second paragraph of 35 U.S.C. 112:

The specification shall conclude with one or more claims particularly pointing out and distinctly claiming the subject matter, which the applicant regards as his invention.

Claims 1-24, 28-31 and 36 are rejected under 35 U.S.C. 112, second paragraph, as being indefinite for failing to particularly point out and distinctly claim the subject matter which applicant regards as the invention.

Claim 1 recites "wherein the HIV envelope protein is adapted to reduce or prevent glycosylation in a mammalian cell." This language is not clear. As written, it seems the HIV protein now has the function of patrolling the target cell and reducing or preventing glycosylation of other proteins.

Claims 7, 9, 11, 14, 19 and 20 recite fusion proteins as A-B-C-D, for example. It is not clear if this is meant to denote the arrangement of the sequences in the 5' to 3' direction.

Claim 16 is drawn to a polynucleotide where a 5' untranslated region between the promoter and the coding sequence comprises exon 1. It is unclear how an untranslated region can code for an exon. If applicants meant to state "wherein a 5' untranslated region including exon1 of the HCMV IE gene is between the HCMV IE promoter and the coding sequences," then the claim should be amended for clarity.

Claim Rejections - 35 USC § 102

The following is a quotation of the appropriate paragraphs of 35 U.S.C. 102 that form the basis for the rejections under this section made in this Office action:

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A person shall be entitled to a patent unless –

(b) the invention was patented or described in a printed publication in this or a foreign country or in public use or on sale in this country, more than one year prior to the date of application for patent in the United States.

Claims 1, 2, 4, 5, 12, 13, 17, 18, 21, 22, 28-31 and 36 are rejected under 35 U.S.C. 102(b) as being anticipated by Nabel et al. (WO 02/32943) as evidenced by Fynan et al. (Proc. Natl. Acad. Sci. USA, 90: 11478-82, 1993).

Nabel et al. discloses various double-stranded DNA vectors comprising sequences that encode an HIV Env that is non-glycosylated (see pages 43, 45 and 46, in particular page 46, lines 13-17) and HIV Env fused to another HIV gene such as Nef (see pages 58-59). The teachings of Nabel et al. include the use of gp120 in the envelope-containing vectors and fusion proteins (see, for example, page 20). The sequences are under the control of a heterologous promoter (see page 23, lines 6-29), and the env sequences are codon optimized for expression in human cells (page 16, line 30 and pages 58-59). Compositions comprising the vectors can be in aqueous solution (page 25, lines 3-6), mixed with a pharmaceutically acceptable adjuvant (page 27, lines 1-15), enclosed in liposome carriers (i.e., particle carriers) (page 25, lines 9-15), and can be used in a prime-boost regimen (page 27, lines 26-27). In addition, Nabel et al. teaches that the compositions can be administered intramuscularly, intraperitoneally, intradermally, subcutaneously, etc. via intradermal delivery devices such as syringes, needle-less injection devices, or microprojectile bombardment gene guns (page 33, lines 5-13). It is well known in the art that gold beads are used as carriers for DNA vaccinations via gene guns or microprojectile bombardment as evidenced by Fynan et al.

Claim Rejections - 35 USC § 103

The following is a quotation of 35 U.S.C. 103(a) which forms the basis for all obviousness rejections set forth in this Office action:

(a) A patent may not be obtained though the invention is not identically disclosed or described as set forth in section 102 of this title, if the differences between the subject matter sought to be patented and the prior art are such that the subject matter as a whole would have been obvious at the time the invention was made to a person having ordinary skill in the art to which said subject matter pertains. Patentability shall not be negated by the manner in which the invention was made.

Claims 15 and 16 are rejected under 35 U.S.C. 103(a) as being unpatentable over Nabel et al. (WO 02/32943) in view of Catchpole et al. (WO 02/36792).

Claims 15 and 16 are drawn to the polynucleotides of the invention linked to the HCMV IE gene promoter and also the 5' untranslated region of the HCMV IE promoter including exon 1 of the HCMV IE gene.

The teachings of Nabel et al. are outlined above under 35 U.S.C. 102(b). Nabel et al. does disclose the use of the CMV promoter (page 31, lines 31-32), but not the use of the HCMV IE gene promoter or the 5' untranslated region of the HCMV IE promoter including exon 1 of the HCMV IE gene.

Catchpole et al. teaches that the use of the HCMV IE gene promoter to drive gene expression is known (page 1, lines 19-33). Catchpole et al. further states that the 5' untranslated region of the HCMV IE promoter including exon 1 of the HCMV IE gene will result in an enhanced level of expression from the HCMV IE promoter (page 2, line 25 to page 4, line 2). Catchpole et al. also teaches that such promoter sequences are good for the expression of HIV antigens (page 5, line 32 to page 7, line 4).

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It would have been obvious to one of ordinary skill in the art to modify the polynucleotides taught by Nabel et al. to use the HCMV IE promoter and 5' sequences from HCMV IE in order to express HIV antigens/genes. One would have been motivated to do so, given the suggestion by Nabel et al. to use the CMV promoter generally to express HIV antigens and the suggestion by Catchpole et al. to use the HCMV IE promoter specifically to drive expression of recombinant proteins and to use the 5' untranslated sequences for enhanced expression of antigens. There would have been a reasonable expectation of success, given the fact that it is well known that the CMV promoter is a strong promoter and that the CMV promoter is commonly used to express heterologous proteins/antigens (see Catchpole et al., page 1, lines 19-33), including HIV antigens. Thus, the invention as a whole was clearly *prima facie* obvious to one of ordinary skill in the art at the time the invention was made.

Claims 23 and 24 are rejected under 35 U.S.C. 103(a) as being unpatentable over Nabel et al. (WO 02/32943) in view of Farina et al. and Roy et al.

The teachings of Nabel et al. are outlined above under 35 U.S.C. 102(b). Nabel et al. does disclose the use of adenovirus as a vector (page 25, lines 16-35), but not a replication-defective adenovirus or Pan 9, 5, 6, or 7.

Farina et al. teaches the use of replication-defective adenovirus C68 to express genes. C68 is another name for Pan 9 as evidenced by Roy et al.

It would have been obvious to one of ordinary skill in the art to modify the polynucleotides taught by Nabel et al. to use the C68 (Pan 9) replication-defective

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adenovirus of Farina et al. One would have been motivated to do so, given the suggestion by Farina et al. that C68 functions as an excellent vaccine for HIV (see Farina et al. page 11612, last paragraph of the discussion) and that humans rarely have neutralizing antibodies to these viruses. There would have been a reasonable expectation of success, given the fact that it is well known to use replication-defective adenoviruses as vaccine vehicles. Thus, the invention as a whole was clearly *prima facie* obvious to one of ordinary skill in the art at the time the invention was made.

Claim 3 is rejected under 35 U.S.C. 103(a) as being unpatentable over Nabel et al. (WO 02/32943) in view of Botarelli et al. (Journal of Immunology, 1991, 147(9):3128-3132).

The teachings of Nabel et al. are outlined above under 35 U.S.C. 102(b). Botarelli et al. teaches that glycosylation residues on gp120 can function as hindering structures that limit antigen recognition by T-lymphocytes (see abstract). The non-glycosylated form of HIV gp120 of Botarelli et al. was produced by removing the signal sequence. Botarelli et al. states that "[t]he lack of signal sequence prevents passage through the secretory pathway and addition of carbohydrates."

It would have been obvious to one of ordinary skill in the art to modify the polynucleotides taught by Nabel et al. to use HIV env sequences lacking a secretory signal sequence to produce non-glycosylated HIV Env. One would have been motivated to do so, given the suggestion by Botarelli et al. that glycosylation residues on gp120 can function as hindering structures that limit antigen recognition by T-

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lymphocytes. There would have been a reasonable expectation of success, given the fact that it is well known that the removal of the secretory signal sequence bypasses the secretory pathway and the addition of carbohydrates, and that others have successfully produced non-glycosylated proteins by removing the secretory signal sequence (see Botarelli et al.). Thus, the invention as a whole was clearly *prima facie* obvious to one of ordinary skill in the art at the time the invention was made.

Claims 1, 2, 4 and 12 are rejected under 35 U.S.C. 103(a) as being unpatentable over Jiang et al. (Chinese Journal of Microbiology and Immunology, 2002, 22(5):482-484)(Abstract only) in view of Botarelli et al. (Journal of Immunology, 1991, 147(9):3128-3132).

Jiang et al. teaches an HIV gp120-gag fusion protein antigen expressed in yeast from the yeast expression vector pHLS1.

Botarelli et al. teaches that glycosylation residues on gp120 can function as hindering structures that limit antigen recognition by T-lymphocytes (see abstract). The non-glycosylated form of HIV gp120 of Botarelli et al. was produced by removing the signal sequence. Botarelli et al. states that "[t]he lack of signal sequence prevents passage through the secretory pathway and addition of carbohydrates."

It would have been obvious to one of ordinary skill in the art to modify the polynucleotides taught by Jiang et al. to use HIV gp120 sequences lacking glycosylation. This can be achieved by removing the secretory signal sequence as taught by Botarelli et al. or by mutating glycosylation signals. One would have been

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motivated to do so, given the suggestion by Botarelli et al. that glycosylation residues on gp120 can function as hindering structures that limit antigen recognition by T-lymphocytes. There would have been a reasonable expectation of success given the fact that it is well known in the art that mutating glycosylation signals and/or removing the secretory signal sequence to bypasses the secretory pathway both result in non-glycosylated forms of the peptide. Thus, the invention as a whole was clearly *prima facie* obvious to one of ordinary skill in the art at the time the invention was made.

Double Patenting

The nonstatutory double patenting rejection is based on a judicially created doctrine grounded in public policy (a policy reflected in the statute) so as to prevent the unjustified or improper timewise extension of the "right to exclude" granted by a patent and to prevent possible harassment by multiple assignees. A nonstatutory obviousness-type double patenting rejection is appropriate where the conflicting claims are not identical, but at least one examined application claim is not patentably distinct from the reference claim(s) because the examined application claim is either anticipated by, or would have been obvious over, the reference claim(s). See, e.g., *In re Berg*, 140 F.3d 1428, 46 USPQ2d 1226 (Fed. Cir. 1998); *In re Goodman*, 11 F.3d 1046, 29 USPQ2d 2010 (Fed. Cir. 1993); *In re Longi*, 759 F.2d 887, 225 USPQ 645 (Fed. Cir. 1985); *In re Van Ornum*, 686 F.2d 937, 214 USPQ 761 (CCPA 1982); *In re Vogel*, 422 F.2d 438, 164 USPQ 619 (CCPA 1970); and *In re Thorington*, 418 F.2d 528, 163 USPQ 644 (CCPA 1969).

A timely filed terminal disclaimer in compliance with 37 CFR 1.321(c) or 1.321(d) may be used to overcome an actual or provisional rejection based on a nonstatutory double patenting ground provided the conflicting application or patent either is shown to be commonly owned with this application, or claims an invention made as a result of activities undertaken within the scope of a joint research agreement.

Effective January 1, 1994, a registered attorney or agent of record may sign a terminal disclaimer. A terminal disclaimer signed by the assignee must fully comply with 37 CFR 3.73(b).

Claims 1-24, 28-31 and 36 are rejected on the ground of nonstatutory obviousness-type double patenting as being unpatentable over claims 1-20, 24-27 and

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32 of copending Application No. 11/734,464. Although the conflicting claims are not identical, they are not patentably distinct from each other because both sets of claims encompass a recombinant polynucleotide molecule comprising HIV envelope sequences fused to at least one HIV nonstructural protein or capsid protein in a vector with a heterologous promoter.

This is a provisional obviousness-type double patenting rejection because the conflicting claims have not in fact been patented.

Claims 4, 6, 7, 10-14 and 19 are rejected on the ground of nonstatutory obviousness-type double patenting as being unpatentable over claims 1-7, 9, 10, 13-16 and 18-21 of U.S. Patent No. 10/490,011. Although the conflicting claims are not identical, they are not patentably distinct from each other because both sets of claims encompass a recombinant polynucleotide molecule comprising gp120 sequences, a heterologous promoter and an other HIV proteins such as Nef and/or Tat in a vector with an enhanced HCMV IE1 promoter. For example, the gp120-RT-Nef-Gag peptide of instant claim 7 would also read on the claims of co-pending application 10/490,011.

This is a provisional obviousness-type double patenting rejection because the conflicting claims have not in fact been patented.

Any inquiry concerning this communication or earlier communications from the examiner should be directed to Nicole E. Kinsey, Ph.D. whose telephone number is

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(571) 272-9943. The examiner can normally be reached on Monday through Friday from 8:00 am to 5:30 pm.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Bruce Campell can be reached on (571) 272-0974. The fax phone number for the organization where this application or proceeding is assigned is 571-273-8300.

Nicole E. Kinsey, Ph.D.
Examiner
Art Unit 1648

/nk/

/Stacy B. Chen/ 9-5-07
Primary Examiner, TC1600